

REMARKS

The independent claims are claims 9 and 10. Claims 1-8 and 16-22 are withdrawn as being directed to a non-elected invention. Review and reconsideration on the merits are requested.

(1) Claims 9 and 12-14 stand rejected under 35 U.S.C. § 102(b) as anticipated by or, in the alternative, under 35 U.S.C. § 103(a) as obvious over U.S. Patent 6,150,426 to Curtin et al.

Because Curtin et al is said to teach copolymers of TFE and a perfluorinated vinyl ether “preferably substantially free of components containing carbonyl bonds,” the Examiner considered that the intensity ratio $[x/y]$ of not higher than 0.05 in IR measurement is met by Curtin et al. In the Advisory Action dated October 6, 2009, the Examiner maintained that “free of components containing carbonyl bonds” means the absence of all functional groups containing a carbon-oxygen double bond, including carboxyl group. The Examiner further noted that Curtin et al contrast their copolymers “free of components containing carbonyl groups” from those of U.S. Patent 4,433,082 to Grot indicating bands at 1740 cm^{-1} corresponding to the presence of carbonyl groups. Regarding this last point, the Examiner commented that “the bands at 1740 cm^{-1} do not apply to the invention of Curtin, but rather show the difference between Curtin and the prior art.”

Claim 9 is directed to a stabilized fluoropolymer represented by formula (II), which stabilized fluoropolymer shows an intensity ratio $[x/y]$ between carboxyl group-due peak $[x]$ and $-\text{CF}_2-$ due peak $[y]$ no higher than 0.05 in IR measurement.

The rejection should be withdrawn because (i) the polymer of Curtin et al necessarily has a carboxyl at its terminal end and therefore does not meet the claimed intensity ratio $[x/y]$ of not higher than 0.05 in IR measurement; and (ii) the description in Curtin et al to the effect that the

solid compositions described therein are “preferably substantially free of components containing carbonyl bonds” says nothing about whether or not the subject polymer has a carboxyl end group. Applicants explain in further detail below and by reference to the Declaration under 37 C.F.R. § 1.132 of Tadashi Ino submitted concurrently herewith.

Curtin et al relates to a process for preparing a composition comprising particles of highly fluorinated ion-exchange polymer having sulfonate functional groups, which process comprises contacting the polymer in a pressurized vessel with an aqueous liquid dispersion medium under conditions which cause the polymer to form particles, cooling the contents of the vessel, and recovering the desired product.

1. The polymer of Curtin et al necessarily has a carboxyl end group

Curtin et al identifies the polymers disclosed in U.S. Patent 3,282,875 to Connolly et al as being preferred polymers for use in their invention. See col. 4, lines 30-32 of Curtin et al. Although U.S. Patent 3,282,875 and Curtin et al do not disclose an end group of a polymer, GB 1,210,794 discloses that the polymer of U.S. Patent 3,282,875 has an unstable carboxyl group. See Examples 1, 6 and 9 in Table I of GB ‘794 employing the copolymer of Example 8 of U.S. Patent 3,282,875 (page 4, lines 96-103). As shown in Table I of GB ‘794, Examples 1, 6 and 9 not subjected to fluorination treatment had a carboxylate content in excess of 500 end groups/ 10^6 C atoms.

U.S. Patent 3,085,083 to Schreyer at col. 2, lines 28-33 explains how carboxylate end-groups in a fluorocarbon polymer are formed. Particularly, carboxylate end-groups in a fluorocarbon polymer are formed when polymerization is initiated through a peroxide catalyst, or when the polymerization is terminated through the formation of a vinyl bond which is subsequently oxidized. Curtin et al does not disclose a polymerization method, but cites to U.S.

Patent 3,282,875 to Connolly at col. 4, lines 31-32. Connolly et al discloses a method of producing fluorocarbon vinyl ether polymers. In polymerization, a peroxide or a redox initiator is used as an initiator, as illustrated at col. 2, lines 55-60. Namely, in the absence of subsequent treatment to remove carboxyl end groups, such carboxyl end groups are necessarily present in the copolymer of Curtin et al derived from the initiator used in polymerization.

Fluorination treatment is needed to reduce the intensity ratio $[x/y]$. See, for example, the unfluorinated polymer of Examples 1 and 3 of the present specification. Prior to fluorination, the polymer of Example 1 had an intensity ratio $[x/y]$ of 0.23 (page 47, lines 23-25), and the polymer of Example 3 prior to fluorination treatment had an intensity ratio $[x/y]$ of 0.08 (page 51, lines 26-29 of the specification). As shown above, the copolymer of Curtin et al necessarily contains carboxyl end groups derived from the initiator used in polymerization, and Curtin et al does not disclose any technique for removing carboxyl end groups. In fact, Curtin et al's investigation was unrelated to the end group of the fluoropolymer.

2. Curtin does not refer to an end group of the polymer

The passage at col. 6, lines 49-65 of Curtin et al cited by the Examiner is reproduced below.

The solid compositions, including the compositions obtained upon drying preferred liquid compositions in accordance with the invention, are preferably substantially free of components containing carbonyl bonds as determined by reflectance infrared spectroscopy. In contrast, reflectance infrared spectroscopy of solids recovered from compositions made by the process of U.S. Pat. No. 4,433,082 to Grot indicate bands at 1740 cm^{-1} corresponding to the presence of carbonyl groups. It is believed that compounds containing carbonyl groups are formed during manufacturing due to the presence of alcohols in the dispersion process. Preferred compositions in accordance with the invention are also free of C-H bonds, i.e., no bands $2800\text{-}3000\text{ cm}^{-1}$ occur in reflectance infrared spectroscopy, unless they are present in the

polymer molecule, e.g., unfluorinated sites or quaternary amine cation associated with the -SO_3^- group.

The band at 1740 cm^{-1} corresponds to carbonyl groups in low molecular weight organic compounds. Particularly, the “carbonyl groups” are identified in the very same passage as carbonyl group-containing compounds formed during manufacturing due to the presence of alcohols in the dispersion process. That is, the “carbonyl groups” mentioned by Curtin et al are present in by-products and have nothing to do with carboxyl end groups of the copolymer.

When a carboxyl group is present at the end of a fluoropolymer, its band shifts to a larger wave number as shown in Table X below. The IR spectrum of a fluoropolymer having carboxyl end groups does not show bands at 1740 cm^{-1} . That is, the bands at 1740 cm^{-1} do not include bands derived from carboxyl end groups of a fluoropolymer.

Table X

	U.S. Patent 3,085,083	U.S. Patent 4,675,380	U.S. Patent 6,689,833
-COF		1883 cm^{-1}	1888 cm^{-1}
-COOH (monomer)	1812 cm^{-1}	1812 cm^{-1}	1813 cm^{-1}
-COOH (dimmer)	1779 cm^{-1}	1773 cm^{-1}	1774 cm^{-1}
$\text{-CO}_2\text{CH}_3$		1795 cm^{-1}	

The shift is due to low electron density at the carbonyl carbon atom by the electron-withdrawing fluorocarbon group.

3. The reflectance infrared spectroscopy indicates very small bands

There is no disclosure in Curtin et al relating to bands corresponding to the carboxyl end groups of a fluoropolymer.

As shown in Graph 1 of the Declaration under 37 C.F.R. § 1.132 of Tadashi Ino submitted herewith, the reflectance infrared spectroscopy indicates very small bands corresponding to the presence of carboxyl end groups of the fluoropolymer. It is easy to miss these bands and, therefore, careful observation is needed to recognize the same.

Because Curtin et al were not interested in the end group of a fluoropolymer, it is not surprising that Curtin et al do not refer to the presence of bands other than those at 1740 cm^{-1} .

The present Applicants have submitted more than adequate rebuttal evidence, demonstrating that the copolymer of Curtin et al necessarily has carboxyl end groups. As noted above and in the Declaration of Mr. Ino, such carboxyl end groups are derived from the initiator used in polymerization. GB '794 and U.S. Patent 3,282,875 to Connolly et al cited by Curtin et al confirm that the copolymer of Curtin et al contains carboxyl end groups and in large amount. Curtin et al does not even mention the copolymer end groups, much less any fluorination treatment or otherwise for converting carboxyl end groups. As explained in Mr. Ino's Declaration, the disclosure "substantially free of components containing carbonyl bonds" of Curtin et al has nothing to do with the polymer end groups.

For the above reasons, it is respectfully submitted that claims 9 and 12-14 are neither anticipated nor obvious over Curtin et al, and withdrawal of the foregoing rejection is respectfully requested.

(2) Claims 10 and 11 stand rejected under 35 U.S.C. § 103(a) as being unpatentable over Curtin et al in view of U.S. Patent 3,085,083 to Schreyer. Curtin et al was cited as

disclosing that preferred compositions are “free of C-H bonds.” Schreyer was cited as teaching -CF₂H end groups. The reason for rejection was that it would have been obvious that “given the end group -CF₂H of Schreyer on a fluoropolymer that is free of C-H bonds as taught by Curtin would result in the claimed end groups of -CF₃.”

Applicants respond as follows.

Claim 10 is directed to a stabilized fluoropolymer represented by general formula (II), wherein in a hydrolyzate of the stabilized fluoropolymer, the number [x] of main chain terminal -CF₃ groups per 1×10^5 main chain carbon atoms of the hydrolyzate is not smaller than 10 (as calculated using an integrated intensity due to main chain terminal -CF₃ groups and an integrated intensity due to -CF₂- adjacent to an ether bond and side chains branched from the main chain in the hydrolyzate).

Turning to the cited prior art, as explained in the Declaration under 37 C.F.R. § 1.132 submitted herewith, the polymer of Curtin et al has a carboxyl end group. Further, there is no description in Curtin et al to the effect that the end group is fluorinated. Thus, the number [X] of main chain terminal -CF₃ groups per 1×10^5 main chain carbon atoms is less than 10.

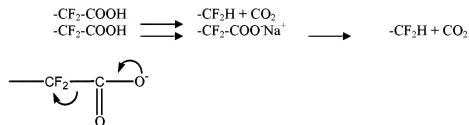
The combination of Curtin et al and Schreyer does not teach the present invention. Particularly, the polymer is necessarily fluorinated to obtain a -CF₃ end group.

However, the end group of the polymer of Curtin et al is a carboxyl group. On the other hand, the end group of the polymer of Schreyer is a -CF₂H group such that the combination of Curtin et al results in a -CF₂H group even if Schreyer teaches that additional fluorinated methyl end groups add stability.

Moreover, neither Curtin et al nor Schreyer et al teach any stabilization method by fluorination. Thus, the combination does not provide a polymer having a -CF₃ end group.

In more detail, Schreyer teaches two techniques to carry out end-capping bridging cols. 3-4, including slurry-capping and vapor-capping. Basically, in Schreyer's technique, the polymer is treated with water at elevated temperatures to form a substantial number of stable $\text{-CF}_2\text{H}$ groups. See col. 2, lines 63-68 of Schreyer. As disclosed at col. 4, lines 10-14, the completion of the reaction, namely, the transformation of substantially all of the carboxylate end-groups to $\text{-CF}_2\text{H}$ end-groups, can be determined readily by infrared techniques.

Importantly, as explained by Mr. Ino, the treatment of Schreyer cannot form -CF_3 end groups.



Because Curtin et al does not teach fluorination treatment, and because the treatment of Schreyer cannot form -CF_3 end groups, the resulting combination can never achieve the invention of claim 10.

The statement in Curtin et al that the fluoropolymer is then “free of C-H bonds” refers to a (per)fluoropolymer having $\text{-CF}_2\text{-}$ but no $\text{-CH}_2\text{-}$ or -CFH- in the main chain and has nothing to do with the terminal groups. That is, the perfluoropolymer of Curtin et al will always have end groups “free of C-H bonds,” unless it is a quaternary amine cation associated with the SO_3 group (which Curtin et al explain is one of the exceptions to “free of C-H bond”).

As explained by Mr. Ino, if there are unfluorinated sites such as VDF unit in the polymer, the polymer indicates bands at $2800\text{-}3000\text{ cm}^{-1}$. The polymer with $\text{-SO}_3\text{NR}_4$ also indicates bands

at 2800-3000 cm^{-1} . However, there is no such band in the IR-profile in Mr. Ino's Declaration because the polymer has neither unfluorinated sites nor quaternary amine cation.

Thus, "free of C-H bonds" does not mean "free of unstable carboxyl end groups."

Withdrawal of the foregoing rejection under 35 U.S.C. § 103(a) is respectfully requested.

(3) Claim 15 stands rejected under 35 U.S.C. § 103(a) as being unpatentable over Curtin et al in view of WO 02/096983 A1.

Applicants rely on the response above with respect to the rejection of claims 9 and 12-14 over Curtin et al alone. Withdrawal of the foregoing rejection under 35 U.S.C. § 103(a) is respectfully requested.

Withdrawal of all rejections and allowance of claims 9-15 is earnestly solicited.

In the event that the Examiner believes that it may be helpful to advance the prosecution of this application, the Examiner is invited to contact the undersigned at the local Washington, D.C. telephone number indicated below.

The USPTO is directed and authorized to charge all required fees, except for the Issue Fee and the Publication Fee, to Deposit Account No. 19-4880. Please also credit any overpayments to said Deposit Account.

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